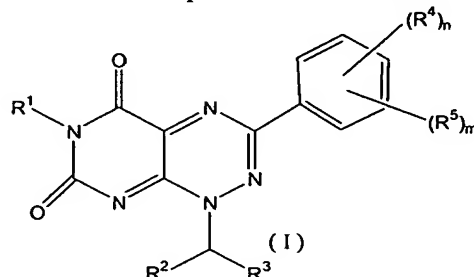


ABSTRACT3-PHENYL ANALOGS OF TOXOFLAVINE AS KINASE INHIBITORS

The present invention concerns the compounds of formula



5

the N-oxide forms, the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, wherein n represents an integer being 0, 1 or 2; m represents an integer being 0 or 1; R¹ represents C₁₋₄alkyl; R² represents C₁₋₄alkyl; R³ represents C₁₋₄alkyl; or R² and R³ taken together with the carbon atom to which they are attached form a C₃₋₈cycloalkyl or Het¹ wherein said C₃₋₈cycloalkyl or Het¹ each independently may optionally be substituted with C₁₋₄alkyloxycarbonyl; R⁴ represents halo or C₁₋₄alkyloxy; R⁵ represents C₁₋₄alkyloxycarbonyl, -O-(mono- or di(C₁₋₄alkyl)aminosulfonyl), C₁₋₄alkyl substituted with one or where possible more substituent being selected from Het³ or NR⁶R⁷, C₁₋₄alkyloxy substituted with one or where possible more substituents being selected from amino, Het⁴ or NR⁸R⁹; R⁶ and R⁷ are each independently selected from hydrogen, C₁₋₄alkyl, C₁₋₄alkyloxyC₁₋₄alkyl, -Het⁵ or C₁₋₄alkyl substituted with one or where possible more substituents being selected from hydroxy, or Het⁵; R⁸ and R⁹ are each independently selected from hydrogen, C₁₋₄alkyl, -Het⁷ or mono- or di(C₁₋₄alkyl)aminosulfonyl; Het³ represents a heterocycle selected from piperidinyl, or piperazinyl wherein said monocyclic heterocycles each independently may optionally be substituted with one, or where possible two or three substituents each independently selected from hydroxy, aminosulfonyl, amino, mono- or di(C₁₋₄alkyl)aminosulfonyl, hydroxyC₁₋₄alkyloxyC₁₋₄alkyl or C₁₋₄alkyloxy; Het⁴ represents a heterocycle selected from morpholinyl, piperidinyl or piperazinyl wherein said monocyclic heterocycles each independently may optionally be substituted with one, or where possible two or three substituents each independently selected from C₁₋₄alkyl, C₁₋₄alkyloxycarbonyl or mono- or di(C₁₋₄alkyl)aminosulfonyl; Het⁵ represents a heterocycle selected from pyridinyl or piperidinyl wherein said monocyclic heterocycles each independently may optionally be substituted with one, or where possible two or three substituents each independently selected from aminosulfonyl, or mono- or di(C₁₋₄alkyl)aminosulfonyl; Het⁷ represents piperidinyl.